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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/526,323	10/03/2005	Richard H Ebright	744-47 PCT/US	6511
23869 7590 04/24/2008 HOFFMANN & BARON, LLP 6900 JERICHO TURNPIKE SYOSSET, NY 11791				
EXAMINER				
HA, JULIE				
ART UNIT		PAPER NUMBER		
1654				
MAIL DATE		DELIVERY MODE		
04/24/2008		PAPER		

**Please find below and/or attached an Office communication concerning this application or proceeding.**

The time period for reply, if any, is set in the attached communication.

### Office Action Summary

**Application No.**

10/526,323

**Applicant(s)**

EBRIGHT, RICHARD H

**Examiner**

JULIE HA

**Art Unit**

1654

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --  
**Period for Reply**

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

**Status**

- 1) ☒ Responsive to communication(s) filed on 19 February 2008.
- 2a) ☒ This action is **FINAL**. 2b) ☐ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

**Disposition of Claims**

- 4) ☒ Claim(s) 78-90, 92-101 and 103-120 is/are pending in the application.
- 4a) Of the above claim(s) 78-83 and 104-120 is/are withdrawn from consideration.
- 5) ☐ Claim(s) \_\_\_\_\_ is/are allowed.
- 6) ☒ Claim(s) 84-90, 92-101 and 103 is/are rejected.
- 7) ☐ Claim(s) \_\_\_\_\_ is/are objected to.
- 8) ☐ Claim(s) \_\_\_\_\_ are subject to restriction and/or election requirement.

**Application Papers**

- 9) ☐ The specification is objected to by the Examiner.
- 10) ☐ The drawing(s) filed on \_\_\_\_\_ is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.  
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).  
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
- 11) ☐ The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

**Priority under 35 U.S.C. § 119**

- 12) ☐ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
- a) ☐ All b) ☐ Some \* c) ☐ None of:
1. ☐ Certified copies of the priority documents have been received.
  2. ☐ Certified copies of the priority documents have been received in Application No. \_\_\_\_\_.
  3. ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

\* See the attached detailed Office action for a list of the certified copies not received.

**Attachment(s)**

- 1) ☐ Notice of References Cited (PTO-892)
- 2) ☐ Notice of Draftsperson's Patent Drawing Review (PTO-948)
- 3) ☐ Information Disclosure Statement(s) (PTO/SB/08)  
Paper No(s)/Mail Date \_\_\_\_\_
- 4) ☐ Interview Summary (PTO-413)  
Paper No(s)/Mail Date \_\_\_\_\_
- 5) ☐ Notice of Informal Patent Application
- 6) ☐ Other: \_\_\_\_\_

### **DETAILED ACTION**

Response to Non-final rejection filed on February 19, 2008 is acknowledged. Claims 78-90, 92-101, 103-120 are pending in this application. Claims 78-83, 104-120 remain withdrawn from further consideration as being drawn to nonelected invention. Claims 84-90, 92-101 and 103 are examined on the merits in this office action.

#### ***Withdrawn Rejection***

1. Rejection under Obviousness Double Patenting is hereby withdrawn due to Applicant's arguments.

#### ***Maintained Rejection***

##### **35 U.S.C. 103**

2. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

3. The factual inquiries set forth in *Graham v. John Deere Co.*, 383 U.S. 1, 148 USPQ 459 (1966), that are applied for establishing a background for determining obviousness under 35 U.S.C. 103(a) are summarized as follows:

1. Determining the scope and contents of the prior art.
2. Ascertaining the differences between the prior art and the claims at issue.
3. Resolving the level of ordinary skill in the pertinent art.
4. Considering objective evidence present in the application indicating obviousness or nonobviousness.

4. Claims 84-90, 92-101 and 103 are rejected under 35 U.S.C. 103(a) as being unpatentable over Delgado et al (Journal of Bacteriology, 2001, 183: 4543-4550) in view of Korzheva et al (Science, 2000, 289: 619-625), Darst et al (PG Pub 2002/0034808) and Woychik (Cell, Feb 2002, 108: 453-463).
5. The instant claims are drawn to method of identifying an agent that inhibits RNA synthesis activity of *E. coli* RNAP by binding to an RNAP secondary channel amino acid sequence. The claims are further drawn to comparing the inhibition by an agent that binds to the secondary channel with inhibition by an agent that binds to a mutant channel or one that binds to fragment of RNAP.
6. With respect to claims 84-89 and 93-100, Delgado et al disclose methods to identify an agent that binds to the secondary channel by determining the binding site of the antibiotic MccJ25 within the intact *E. coli* RNAP. Delgado et al further disclose that such methods involve contacting the RNAP with the antibiotic and detecting the inhibition of RNA synthesis in presence of MccJ25 via a transcription assays (see kinetics of binding, Materials and Methods). Delgado et al also disclose a mutant residing in the homology block G of the  $\beta'$  subunit (see abstract) of RNAP and comparative methods employing such a mutant in presence of the antibiotic versus the intact polymerase in presence of the antibiotic to identify the presence or absence of binding. Resistance to the antibiotic was indicative of the antibiotics ability to bind to RNAP in the absence of mutation. Delgado et al disclose that the affected mutation is conserved in all prokaryotic homologues examined (Figure 2), which is good evidence to indicate that the region that contains the mutation is part of the catalytic center of the enzyme, and

that which is responsible for the transcription properties of RNAP. The difference between the reference and the instant claims is that the reference does not explicitly recite binding of the agent to the RNAP secondary channel.

7. However, Korzheva disclose a model based on a bacterial x-ray crystal structure (p. 620, first paragraph) known in the art wherein the secondary channel is responsible for the diffusion of incoming nucleotide substrates into the active site (Figure 3) to overcome access to the main channel which is blocked by the nucleic acid framework.

8. Therefore, it would have been obvious to one of ordinary skill in the art to use the method of identifying an agent that binds to a specific domain of RNAP to inhibit RNA synthesis, and to combine the method with agents with potential to bind to the secondary channel as modeled by Korzheva for the known and expected result of providing a means recognized in the art to identify regions in RNAP that contribute to sensitivity towards agents with inhibition of RNA synthesis activity for the development of a new antibiotic.

### ***Response to Applicant's Arguments***

9. Applicant argues that "the novelty of the instant invention is the discovery that amino acids 736-747 and 779-781 of the RNAP  $\beta'$  subunit are a useful target for compounds that block DNA transcription. This contrasts starkly with the mutation discovered by Delgado in residue 931 of the RNAP  $\beta'$  subunit...the distinct locations of the modifications to the RNAP  $\beta'$  subunit is patentably significant. It should be apparent to a skilled person in the field of molecular biology that different mutations and different

locations on a protein or DNA segment have different function." Applicant further argues that "the present Applicants have identified a specific region of interest in the secondary channel only, and specifically enumerated residues, that are clearly stated in the currently amended claims. The subunit that we are claiming is not discussed in any of the cited references." Furthermore, Applicant argues that "The "wherein" clause in claim 84 serves to distinguish the present invention from Delgado, by explicitly limiting the amino acids of use in the present method to 736-747, and 779-781 of the  $\beta'$  subunit of RNAP...claim 84 is limited to sequences corresponding to, and alignable with, amino acid residues 736-747 and 779-781. This is not a suggested or optional limitation. It is a firm limitation that clearly places the sequences of interest outside the scope of residue 931 studied by Delgado."

10. Applicant's arguments have been considered but have not been found persuasive because Delgado et al disclose the  $\beta'$  subunit of *E. coli* RNAP, and the mutation in this strain was cloned by in vivo recombination into an rpoC+ plasmid. The presence of the recombinant plasmid conferred complete resistance to otherwise sensitive strains. Delgado et al further disclose that these results, along with the observation that MccJ25 inhibits in vivo and in vitro RNA synthesis, provide convincing evidence that RNAP is the target for MccJ25 action (see p. 4543, right column, 1<sup>st</sup> paragraph). Furthermore, the reference discloses that alignment of prokaryotic and eukaryotic RNAP sequences has defined segments of substantial sequence conservation...For  $\beta'$  subunit of prokaryotic RNAP, eight segments with a particularly high degree of conservation have been designated A through H (see p. 4547, left

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column, 2<sup>nd</sup> paragraph and Fig. 2). Applicant is reminded that the amendment to the claims comprises a "wherein clause", which does not carry a patentably weight. The MPEP states the following: Language that suggests or makes optional but does not require steps to be performed or does not limit a claim to a particular structure does not limit the scope of a claim or claim limitation" (see MPEP 2106). The claims are drawn to a method of not to structure. The "wherein" clause of claim 84 limits the claim to a particular structure, not to steps to be performed. Therefore, in this instance, the "wherein" clause carries no patentable weight. The claims are drawn to a method of identifying an agent that binds to a bacterial RNAP homologous secondary channel amino acid, and the residues are in the  $\beta'$  subunit of RNAP of *E. coli*. Since the primary reference teaches the  $\beta'$  subunit of RNAP of *E. coli*, an agent that binds to the  $\beta'$  subunit of RNAP of *E. coli* would necessarily bind to any amino acid residue of  $\beta'$  subunit of RNAP of *E. coli*. The reference teaches the 8 segments of high degree of conservation, and segment F comprises the amino acid residues 700-800. Therefore, since the reference teaches the  $\beta'$  subunit of RNAP of *E. coli*, an agent that binds to the  $\beta'$  subunit of RNAP of *E. coli* would necessarily bind to segment comprising residues 700-800. Furthermore, Korzheva et al teach that the secondary channel is responsible for the diffusion of incoming nucleotide substrate into the active site of RNAP (see abstract and Figure 3), to overcome access to the main channel which is blocked by the nucleic acid framework. Therefore, the claims are unpatentable over the prior arts of record.

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11. With respect to claims 90, 92, 101 and 103, Delgado et al teachings are as discussed above. Further, Delgado disclose that the same comparative methods to detect activity in mutant bacterial RNAP can be applied to eukaryotes. The use of these methods resulted in locating the position of the mutant in yeast (eukaryote). This mutant as disclosed corresponds exactly to the above-mentioned mutant in bacterial RNAP that confers resistance to the antibiotic MccJ25. The difference between the reference and the instant claims is that the reference does not explicitly recite a derivative of human RNA polymerase.

12. However, Darst et al disclose methods of identifying an agent for use as inhibitors of eukaryotic RNAP polymerase (see claim 11). Further, Darst et al disclose that not all residues in the  $\beta'$  subunit of prokaryotic RNAP are conserved in eukaryotic RNAP, pointing to the variable roles of the residues with respect to assembly and/or catalysis within the same subunit (see paragraphs [0176] and [0179]).

13. Woychik et al disclose that human and yeast RNAP II share a much higher level of sequence identity at both the surface and core positions (page 457, paragraph 3). However, there is no significant conservation of surface residues between yeast RNAP II and bacterial RNAP (page 457, paragraph 4).

14. Therefore, it would have been obvious to one of ordinary skill in the art at the time of invention to use the method of identifying an agent that binds to a specific domain of prokaryotic RNAP with eukaryotic RNAP (human or yeast) for the known an expected result of providing a means recognized in the art to identify the basis of differential specificity within species and how that specificity contributes to sensitivity



towards agents capable of inhibiting RNA synthesis activity. Furthermore, it would have been obvious to one of ordinary skill in the art at the time of invention to use the method of identifying an agent that binds to the bacterial RNAP secondary channel wherein agents other than MccJ25 are tested against MccJ25 as a control for the known and expected result of providing a means recognized in the art to compare the binding and inhibition properties of agents against a reference antibiotic known in the art to inhibit RNA synthesis by binding to RNAP.

***Response to Applicant's Arguments***

15. Applicant argues that "Applicants have identified a specific location of the RNAP  $\beta'$  subunit that is nearly invariant among bacterial RNAP, but is significantly different from eukaryotic RNAP...this invariant region is not disclosed in any of the cited art. Furthermore, the inventors have discovered that blocking the RNAP secondary channel with a small molecule prevents uptake of NTPs by RNAP and therefore inhibits transcription." Applicant further argues "the Woychik and Korzheva references are not directed to the inhibition of RNAP mediated transcription. Darst discloses a method for identifying agents that inhibit RNAP activity, but there is no suggestion in Darst to combine his method with the sequence locations identified in the instant application." Furthermore, Applicant argues that "Darst teaches away from any distinction between bacterial and eukaryotic RNAP with the assertion that there is high degree of sequence homology between prokaryotes and eukaryotes." Regarding the KSR analysis, Applicant argues that "the  $\beta'$  subunit of bacterial RNAP has at least 1500 amino acid

residues, and even though it was known that MccJ25 targets the  $\beta'$  subunit of RNAP, substantial experimentation would be required to find other relevant target residues."

16. Applicant's arguments have been considered but have not been found persuasive because, as described supra. Furthermore, Darst et al disclose methods of identifying an agent for use as inhibitors of eukaryotic RNAP, and that not all residues in the  $\beta'$  subunit of prokaryotic RNAP are conserved in eukaryotic RNA; Delgado et al disclose that alignment of prokaryotic and eukaryotic RNAP sequences has defined segments of substantial sequence conservation, which may represent domains with conserved functions. Woychik et al disclose that human and yeast RNAP II share a much higher level of sequence identity at both the surface and core positions. However, there is no significant conservation of surface residues between yeast RNAP II and bacterial RNAP. Since human and yeast share a much higher level of sequence identity, and Darst et al disclose the methods of identifying an agent for use as inhibitors of eukaryotic RNAP, it would have been obvious to one of ordinary skill in the art to try human RNAP species to identify if the agents would work as inhibitors on human RNAP as well as eukaryotic (yeast) RNAP. Furthermore, it has been held that under KSR that "obvious to try" may be an appropriate test under 103. The Supreme Court stated in KSR, When there is motivation "to solve a problem and there are a finite number of identified, predictable solutions, a person of ordinary skill has good reason to pursue the known options within his or her technical grasp. If this leads to anticipated success, it is likely the product not of innovation but of ordinary skill and common sense. In that instance the fact that a combination was obvious to try might show that it was obvious

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under § 103." *KSR Int'l Co. v. Teleflex Inc.*, 127 S. Ct. 1727, \_\_\_, 82 USPQ2d 1385, 1397 (2007). The "problem" facing those in the art was trying to understand the mechanism of action of MccJ25 (peptide antibiotic) and there were a limited number of methodologies available to do so, for example identifying the mutation causing resistance to MccJ25, and trying to identify the mutants affected in the target of the antibiotics. As described supra, Delgado teaches that MccJ25 RNA synthesis both in vivo and in vitro, and these results point to the RNA polymerase as the target of microcin action (see abstract, and p. 4543, right column, 1<sup>st</sup> paragraph). The skilled artisan would have had reason to try locating the sites of microcin action of the RNAP and comparing the MccJ25 mutants to MccJ25 wild-type, with the reasonable expectation that at least one would be successful. Delgado et al disclose  $\beta'$  subunit of the RNAP from *E. coli*, and the agent that binds to the subunit would bind to RNAP. The reference teaches the 8 segments of high degree of conservation, and segment F comprises the amino acid residues 700-800. Therefore, since the reference teaches the  $\beta'$  subunit of RNAP of *E. coli*, an agent that binds to the  $\beta'$  subunit of RNAP of *E. coli* would necessarily bind to segment comprising residues 700-800. Applicant has not shown that the agent would not necessarily bind to segments A through H disclosed in Delgado reference. Further, Darst et al disclose that not all residues in the  $\beta'$  subunit of prokaryotic RNAP are conserved in the eukaryotic RNAP, and Woychik et al disclose that yeast RNAP II share a much higher level of sequence identity at both the surface and core positions with human RNAP, it would have been obvious to try examining those residues both within the surface and core positions and outside the surface and core positions of the

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sequences. The person of ordinary skill in the art would have examined the highly conserved regions, since it is well known in the art that there is a nexus between the conserved region and the binding properties of the protein. Since the segment of A through H is disclosed to be conserved in the  $\beta'$  subunit of E. coli, one of ordinary skill in the art would have been motivated to examine the conserved regions for binding properties. Thus, examining other putative target sites outside the region and comparing the binding to MccJ25 as a control is a "the product not of innovation but of ordinary skill and common sense," leading to the conclusion that invention is not patentable as it would have been obvious.

**35 U.S.C. 112, 1<sup>st</sup>**

17. The following is a quotation of the first paragraph of 35 U.S.C. 112:

The specification shall contain a written description of the invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the same and shall set forth the best mode contemplated by the inventor of carrying out his invention.

18. Claims 86-89, 95 and 97-99 are rejected under 35 U.S.C. 112, first paragraph, as failing to comply with the written description requirement. The MPEP states that the purpose of the written description requirement is to ensure that the inventor had possession, as of the filing date of the application, of the specific subject matter later claimed by him. The courts have stated:

"To fulfill the written description requirement, a patent specification must describe an invention and do so in sufficient detail that one skilled in the art can clearly conclude that "the inventor invented the claimed invention." Lockwood v. American Airlines, Inc., 107 F.3d 1565, 1572, 41 USPQ2d 1961, 1966 (1997); In re Gosteli, 872 F.2d 1008, 1012, 10 USPQ2d 1614, 1618 (Fed. Cir. 1989) (" [T]he description must clearly allow

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persons of ordinary skill in the art to recognize that [the inventor] invented what is claimed."). Thus, an applicant complies with the written description requirement "by describing the invention, with all its claimed limitations, not that which makes it obvious," and by using "such descriptive means as words, structures, figures, diagrams, formulas, etc., that set forth the claimed invention." Lockwood, 107 F.3d at 1572, 41 USPQ2d at 1966. Regents of the University of California v. Eli Lilly & Co., 43 USPQ2d 1398.

19. The MPEP lists factors that can be used to determine if sufficient evidence of possession has been furnished in the disclosure of the Application. These include "level of skill and knowledge in the art, partial structure, physical and/or chemical properties, functional characteristics alone or coupled with a known or disclosed correlation between structure and function, and the method of making the claimed invention. Disclosure of any combination of such identifying characteristics that distinguish the claimed invention from other materials and would lead one of skill in the art to the conclusion that the applicant was in possession of the claimed species is sufficient." MPEP 2163.

20. Further, for a broad generic claim, the specification must provide adequate written description to identify the genus of the claim. In Regents of the University of California v. Eli Lilly & Co., the court stated:

"A written description of an invention involving a chemical genus, like a description of a chemical species, 'requires a precise definition, such as by structure, formula, [or] chemical name,' of the claimed subject matter sufficient to distinguish it from other materials. Fiers, 984 F.2d at 1171, 25 USPQ2d at 1606; In re Smythe, 480 F.2d 1376, 1383, 178 USPQ 279, 284-85 (CCPA 1973) ("In other cases, particularly but not necessarily, chemical cases, where there is unpredictability in performance of certain species or subcombinations other than those specifically enumerated, one skilled in the art may be found not to have been placed in possession of a genus. . . ."). Regents of the University of California v. Eli Lilly & Co., 43 USPQ2d 1398.

21. The MPEP further states that if a biomolecule is described only by a functional characteristic, without any disclosed correlation between function and structure of the sequence, it is "not sufficient characteristic for written description purposes, even when

accompanied by a method of obtaining the claimed sequence." MPEP 2163. The MPEP does state that for generic claim the genus can be adequately described if the disclosure presents a sufficient number of representative species that encompass the genus. MPEP 2163. If the genus has a substantial variance, the disclosure must describe a sufficient variety of species to reflect the variation within that genus. See MPEP 2163. Although the MPEP does not define what constitute a sufficient number of representative, the Courts have indicated what do not constitute a representative number species to adequately describe a broad generic. In Gostelli, the Court determined that the disclosure of two chemical compounds within a subgenus did not describe that subgenus. In re Gostelli, 872 F.2d at 1012, 10 USPQ2d at 1618.

22. In the instant case, the claims are drawn to a method for identifying an agent that binds to a bacterial RNAP homologous secondary channel amino acid sequence in a first entity, wherein the first entity is selected from the group consisting of a derivative of *E. coli* RNAP and a derivative of *B. subtilis* RNAP, wherein the derivatives contain a bacterial RNAP homologous secondary channel amino acid sequence having at least one substitution, insertion, or deletion of amino acid residues corresponding to, an alignable with, amino acid residues 736-747 and 779-781 of the  $\beta'$  subunit of *E. coli* RNAP or amino acid residues 740-751 and 783-785 of the  $\beta'$  subunit of *B. subtilis* RNAP, respectively. The generic statements derivatives of *E. coli* and RNAP and *B. subtilis* RNAP, and secondary channel amino acid sequence having at least one substitution, insertion, or deletion of amino acid residues do not provide ample written description for the compounds since the claims do not describe a single structural

feature. The specification does not clearly define or provide examples of what qualify as compounds of the claimed invention.

23. As stated earlier, the MPEP states that written description for a genus can be achieved by a representative number of species within a broad generic. It is unquestionable claim 85 is broad generics with respect all possible compounds encompassed by the claims. The possible structural variations are limitless to any class of amino acids or amino acid mimetics that can be substituted or inserted, and make up the class of sequences and derivatives of *E. coli* and RNAP and *B. subtilis* RNAP. It must not be forgotten that the MPEP states that if a peptide is described only by a functional characteristic, without any disclosed correlation between function and structure of the sequence, it is "not sufficient characteristic for written description purposes, even when accompanied by a method of obtaining the claimed sequence." MPEP 2163. Here, though the claims may recite some functional characteristics, the claims lack written description because there is no disclosure of a correlation between function and structure of the compounds beyond compounds disclosed in the examples in the specification. Moreover, the specification lack sufficient variety of species to reflect this variance in the genus since the specification does not provide any examples of derivatives and variants. The specification is void of organic molecules that functions as a peptide-like molecule that qualify for the functional characteristics claimed as a peptide or a peptide-like molecule or other peptidic molecules and other synthetic peptide or peptide-like molecule that can function as amino acids and amino acid like substituents.

24. The specification is limited to the amino acid substitutions at positions 424, 428, 429, 430, 454, 469, 493, 498, 503, 504, 508, 680, 684, 732, 733, 734, 735, 736, 738, 744, 748, 775, 776, 777, 778, 779, 780, 782, 783, 784, 785, 786, 788, 789, 790, 869, 922, 926, 927, 930, 931, 932, 933, 1136, 1137, 1240, 1241, 1244, 1247 and 1248. The working example describes that random mutagenesis was performed by error-prone PCR amplification, and the mutagenesis procedure yields all possible transition and transversion substitutions (see paragraph [0142]). The specification does not describe any deletion or insertion of amino acid residues. Description of single amino acid substitution at positions 424, 428, 429, 430, 454, 469, 493, 498, 503, 504, 508, 680, 684, 732, 733, 734, 735, 736, 738, 744, 748, 775, 776, 777, 778, 779, 780, 782, 783, 784, 785, 786, 788, 789, 790, 869, 922, 926, 927, 930, 931, 932, 933, 1136, 1137, 1240, 1241, 1244, 1247 and 1248 is not sufficient to encompass numerous other peptide sequences that belong to the same genus. For example, the claims are drawn to amino acid sequence having at least one substitution, insertion, or deletion of amino acid residues corresponding to, and alignable with, amino acid residues 736-747 and 779-781 ( $\beta'$  subunit of *E. coli*) and 740-751 and 783-785 ( $\beta'$  subunit of *B. subtilis*). This implies that there are 12 different amino acid positions and 3 different amino acid positions for substitutions, insertion, or deletion for both subunits of the species. That means that there are at least  $12 \times 20 = 240$  and  $3 \times 20 = 60$ , respectively, different possibilities of substitutions, deletions and insertions. Furthermore, there are non-natural amino acids such as D-amino acids,  $\alpha$ -amino acids,  $\beta$ -amino acids and  $\epsilon$ -amino acids, that would further increase the number of substitution, insertion and deletions of



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amino acid residues. Having at least one insertion would increase the numbers of possibilities even further. There are varying lengths, varying amino acid compositions, and numerous distinct qualities that make up the genus. For example, the specification only describes single amino acid substitutions of 736, 738, 744, 779 and 780 (*E. coli*) and 744, 748, 783, 784 and 785. However, there are no examples of deletions and insertions. There is not sufficient amount of examples provided to encompass the numerous characteristics of the whole genus claimed.

25. The description requirement of the patent statute requires a description of an invention, not an indication of a result that one might achieve if one made that invention.

See In re Wilder, 736 F.2d 1516, 1521, 222 USPQ 369, 372-73 (Fed. Cir. 1984)

(affirming rejection because the specification does "little more than outlin[e] goals appellants hope the claimed invention achieves and the problems the invention will hopefully ameliorate"). Accordingly, it is deemed that the specification fails to provide adequate written description for the genus of the claims and does not reasonably convey to one skilled in the relevant art that the inventor(s), at the time the application was filed, had possession of the entire scope of the claimed invention.

26. In the previous office action, the rejection cited that to provide evidence of possession of a claimed genus, the specification must provide sufficient distinguishing identifying characteristics of the genus. The factors to be considered include disclosure of complete or partial structure, physical and/or chemical properties, functional characteristics, structure/function correlating, methods of making the claimed product, or any combination thereof. In this case, claims 86, 89, 95 and 99 are directed to

identifying an agent that binds to RNAP secondary channel and derivatives of RNAP secondary channel having at least one substitution, insertion or deletion. Further, the claims are drawn to methods of identifying an agent that binds to derivative of an eukaryotic RNAP derivative when compared with an agent that binds to a bacterial RNAP secondary channel. While the specification has adequate written description of the RNAP secondary channel, there is no disclosure on the structural limitations of the genus represented by the derivatives of RNAP secondary channel and the genus represented by eukaryotic RNAP. Further, there is no disclosure of the activity of the above-mentioned derivatives, nor any method to analyze the activity of the derivatives. There is no description of the identifying characteristics for recognizing that an agent will inhibit the activity of the derivative of RNAP. One skilled in the art would conclude that the disclosure of intact RNAP secondary channel or eukaryotic RNAP is not representative of the undefined genus of derivatives recited in the claims. Accordingly, in the absence of sufficient recitation of distinguishing identifying characteristics, the specification does not provide adequate written description of the claimed genus. Therefore, the inventor, at the time the application was filed was not in possession of the broad genus comprising "derivatives of bacterial RNAP secondary channel" and "derivatives of eukaryotic RNAP" needed to practice the claimed invention.

### ***Response to Applicant's Arguments***

27. Applicant argues that "disclosure in the specification (p. 14, line 23 to p. 15 line 9, Example 1, Table 2 and Table 5) make clear that the examples of the invention are

clearly defined and described." Applicant further argues that "Tables 2 and 5 show a substantial number of actual substitutions and one deletion to support the claimed invention. Accordingly, Applicants believe the suggestion that Table 2 and 5 do not support the alleged lack of written description is erroneous."

28. Applicant's arguments have been fully considered but have not been found persuasive because Table 2 and 5 only shows some of the possible substitutions, and as Applicant has indicated, Table 5 shows only 1 deletion at position 735. Furthermore, Applicant has indicated that "the  $\beta'$  subunit of bacterial RNAP has at least 1500 amino acid residues in response to 103(a) rejection (see p. 6 of Applicant's remarks). Table 5 shows about 80 substitutions and 1 deletion. Further, Table 5 shows 4 substitutions in the range of residues 736-747 and 4 substitutions in the range of residues 779-781. This is not enough to encompass the different variety of substitutions, insertions and deletions in the RNAP homologous secondary channel that is alignable with amino acid residues 736-747 and 779-781. Furthermore, the claims recite "at least one substitution, insertion or deletion". The specification does not provide ample examples of all possible substitutions, insertions or deletions allowed in the 1500 amino acid residues that align with the amino acid residues 736-747 and 779-781. There may be multiple numbers of insertions, deletions, and substitutions in the RNAP secondary channel amino acid sequence that is alignable to residues 736-747 and 779-781 of  $\beta'$  subunit of RNAP from *E. coli*. Furthermore, since the  $\beta'$  subunit of RNAP has at least 1500 amino acid residues, there are vast numbers of possible homologs that would align with the recited residues of *E. coli*. Accordingly, it is deemed that the specification fails to provide

adequate written description for the genus of the claims and does not reasonably convey to one skilled in the relevant art that the inventor(s), at the time the application was filed, had possession of the entire scope of the claimed invention.

**35 U.S.C. 112, 2<sup>nd</sup>**

29. The following is a quotation of the second paragraph of 35 U.S.C. 112:

The specification shall conclude with one or more claims particularly pointing out and distinctly claiming the subject matter which the applicant regards as his invention.

30. Claim 86, recites the limitation "amino acid residues 740-751 and 783-785 of the  $\beta'$  subunit of *Bacillus subtilis* RNAP" in 6<sup>th</sup> and 7<sup>th</sup> line of the claim. There is insufficient antecedent basis for this limitation in the claim. The base claim 84 only recites that the bacterial RNAP homologous secondary channel amino acid sequence corresponds to, and is alignable with, amino acid residues 736-747 and 779-781 of the  $\beta'$  subunit of RNAP from *E. coli*. The base claims does not recite  $\beta'$  subunit of *Bacillus subtilis* RNAP, therefore, claim 86 lacks antecedent basis.

31. Claim 89 recites the limitation "a derivative of *Bacillus subtilis* RNAP" in 2<sup>nd</sup> and 3<sup>rd</sup> line of the claim and "amino acid residues 740-751 and 783-785 of the  $\beta'$  subunit of *Bacillus subtilis* RNAP" in 6<sup>th</sup> and 7<sup>th</sup> line of the claim. There is insufficient antecedent basis for this limitation in the claim. The base claim 84 only recites that the bacterial RNAP homologous secondary channel amino acid sequence corresponds to, and is alignable with, amino acid residues 736-747 and 779-781 of the  $\beta'$  subunit of RNAP from *E. coli*. Furthermore, claim 87, which claim 89 is dependent from, only recites the

$\beta'$  subunit of RNAP from *E. coli*. The base claims does not recite  $\beta'$  subunit of *Bacillus subtilis* RNAP, therefore, claim 89 lacks antecedent basis.

32. Claim 95 recites the limitation "a derivative of *Bacillus subtilis* RNAP" in 2<sup>nd</sup> and 3<sup>rd</sup> line of the claim and "amino acid residues 740-751 and 783-785 of the  $\beta'$  subunit of *Bacillus subtilis* RNAP" in 6<sup>th</sup> and 7<sup>th</sup> line of the claim. There is insufficient antecedent basis for this limitation in the claim. The base claim 84 only recites that the bacterial RNAP homologous secondary channel amino acid sequence corresponds to, and is alignable with, amino acid residues 736-747 and 779-781 of the  $\beta'$  subunit of RNAP from *E. coli*. Furthermore, claim 93, which claim 95 is dependent from, only recites the  $\beta'$  subunit of RNAP from *E. coli*. The base claims does not recite  $\beta'$  subunit of *Bacillus subtilis* RNAP, therefore, claim 95 lacks antecedent basis.

33. Claim 99 recites the limitation "a derivative of *Bacillus subtilis* RNAP" in 2<sup>nd</sup> and 3<sup>rd</sup> line of the claim and "amino acid residues 740-751 and 783-785 of the  $\beta'$  subunit of *Bacillus subtilis* RNAP" in 6<sup>th</sup> and 7<sup>th</sup> line of the claim. There is insufficient antecedent basis for this limitation in the claim. The base claim 84 only recites that the bacterial RNAP homologous secondary channel amino acid sequence corresponds to, and is alignable with, amino acid residues 736-747 and 779-781 of the  $\beta'$  subunit of RNAP from *E. coli*. Furthermore, claim 97, which claim 99 is dependent from, only recites the  $\beta'$  subunit of RNAP from *E. coli*. The base claims does not recite  $\beta'$  subunit of *Bacillus subtilis* RNAP, therefore, claim 95 lacks antecedent basis.

34. Claims 86, 89, 95 and 99 are rejected under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject

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matter which applicant regards as the invention. The claims recite "derivatives of *E. coli* and RNAP and *B. subtilis* RNAP". It is unclear what type of modifications and changes would constitute derivatives of *E. coli* and RNAP and *B. subtilis* RNAP. For example, there are physical, enzymatic or chemical changes that can affect changes in the structure of compounds that may constitute a derivative of *E. coli* and RNAP and *B. subtilis* RNAP.

### ***Response to Applicant's Arguments***

35. Applicant argues that "the reference to *Bacillus subtilis* in claim 86 is not dependent on claim 84, the base claim. The reference to claim 84 is only to the "first entity" in claim 84. Claim 86 provides the additional limitation that the first entity from claim 84 is selected from *E. coli* or *B. subtilis*, and includes additional detail on substitutions, insertions, or deletions at the enumerated ranges of RNAP of those two bacteria." Applicant argues the same arguments regarding lack of antecedent basis on claims 89, 95 and 99.

36. Applicant's arguments have been fully considered but have not been found persuasive because base claim 84 (and 87, 93 and 97) is drawn to bacterial RNAP from *E. coli*. Dependent claim must further limit the base claim not broaden the scope. Claim 84 (87, 93 and 97) recites "...bacterial RNAP homologous secondary channel amino acid sequence in a first entity, comprising preparing a reaction solution including the agent to be tested and a first entity including a bacterial RNAP homologous secondary channel amino acid sequence...wherein said bacterial RNAP homologous secondary

channel amino acid sequence corresponds to, and is alignable with, amino acid residues 736-747 and 779-781 of  $\beta'$  subunit of RNAP from *E. coli*." The base claim clearly indicates that the bacterial RNAP homologous secondary channel amino acid is from *E. coli* only. However, claim 86 (89, 95 and 99) recites *E. coli* or *B. subtilis*. Again, claim 84 (87, 93, 97) is only drawn to *E. coli*, therefore, by reciting *B. subtilis*, claim 86 (89, 95, 99) is broader than claim 84 (87, 93, 97). Thus, claims 86, 89, 95 and 99 lack antecedent basis.

### **Conclusion**

37. **THIS ACTION IS MADE FINAL.** Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a). No claims are allowed.

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the mailing date of this final action.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to JULIE HA whose telephone number is (571)272-5982. The examiner can normally be reached on Mon-Thurs, 5:30 AM to 4:00 PM.

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If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Cecilia Tsang can be reached on 571-272-0562. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

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Examiner, Art Unit 1654

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Primary Examiner, Art Unit 1654